(MIRA 16:4)

LYUTAYA, M.D.; SAMSONOV, G.V. Preparation and properties of lanthanum nitride. Ukr.khim.zhur. 29: 251-255 163.

> 1. Institut metallokeramiki i spetsial'nykh splavov AN UkrSSR. (Lanthanum compounds) (Nitrides)

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

SAMSONOV, G.V.; LYUTAYA, M.D.; NESHPOR, V.S.

Preparation and physicochemical properties of scandium nitride. Zhur. prikl. khim. 36 no.10:2108-2115 0 '63. (MIRA 17:1)

1. Institut metallokeramiki i spetsial'nykh splavov AN UkrSSR.

5(2) AUTHORS:

Tananayev, I. V., Lyutaya, M. D.

SOV/78-4-1-20/48

TITLE:

I. On the Mixed Hexanitritonickelates of Lanthanum and Potassium (I. O smeshannykh geksanitronikeleatakh lantana i kaliya)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 1, pp 97-102

(USSR)

ABSTRACT:

S lubility in the system $La(NO_3)_3 - K_4 \left[Ni(NO_2)_6\right] - H_2O$ was investigated at 25°. The solid phases separated out were analyzed and the thermograms of these compounds drawn. The solubility curves indicate the gradual formation of three solid phases with a rise of K_4 Ni(NO₂) 6 content. The

following solid phases are formed: K6La2 Ni(NO2)6 3; $K_{21}La_5[Ni(NO_2)_6]_9.H_2O; K_5La[Ni(NO_2)_6]_2.H_2O.$ The individuality

of these compounds was proved by their thermograms. The thermograms of $K_6La_2[Ni(NO_2)_6]_3$ show an endothermic effect

within the temperature range 220-265°. Thereby the color of the salts changes from brown to black. The thermograms of

Card 1/3

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SOV/78-4-1-20/48 I. On the Mixed Hexanitritonickelates of Lanthanum and Potassium

 $K_{21}La_5 \begin{bmatrix} Ni(NO_2)_6 \end{bmatrix} 9 \cdot H_{20}$ show two endothermic effects. The first effect at 1300 indicates the dehydration of the salt. The second one at 230-2700 indicates the decomposition of the salt. The thermogram of $K_5La \begin{bmatrix} Ni(NO_2)_6 \end{bmatrix} 2 \cdot H_{20}$ shows an endothermic effect at 1300 indicating the dehydration of the salt and an endothermic effect at 230-2500 indicating the decomposition of the salt. The solubility of $K_5La \begin{bmatrix} Ni(NO_2)_6 \end{bmatrix} 2 \cdot H_{20} \end{bmatrix}$

in KNO $_2$ solutions (1-7 mol/l) was investigated. It was found that at the same time salting out takes place whereby the solid initial phase is changed to $K_6La_2[Ni(NO_2)_6]$ 3. Rare earths can be separated by KNO $_2$ solutions by fractional crystallization of their mixed hexanitritonickelates. There are 6 figures, 4 tables, and 2 references.

Card 2/3

5(2) AUTHORS:

Tananayev, I. V., Lyutaya, M. D.

SOV/78-4-1-21/48

TITLE:

II. On Mixed Hexanitritonickelates of Praseodymium and

Neodymium (II. O smeshannykh geksanitronikeleatakh prazeodima

i neodima)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 1, pr 103-109

(USSR)

ABSTRACT:

Solubility in the systems $Pr(NO_3)_3 - K_4[Ni(NO_2)_6] - H_2O$ and

 $Nd(NO_3)_3-K_4[Ni(NO_2)_6]-H_2O$ was investigated at 25°C. In the

first system the phases $K_{21}Pr_{5}[Ni(NO_{2})_{6}]9.H_{2}O$ and

K₅Pr[Ni(NO₂)₆]₂.H₂O are gradually formed. The second system

also shows the gradual formation of two solid phases: $K_{21}Nd_5[Ni(NO_2)_6]_9$ - H_2O and $K_5Nd[Ni(NO_2)_6]_2$ - H_2O . The thermograms

were drawn and are shown in the figures 3, 4, and 8, 9. The solubility of K_5 Pr Ni(NO₂)6 2. H_2 0 and K_5 Nd Ni(NO₂)6 2. H_2 0 in

KNO, solutions (1-7 mol/1) was investigated. It was found that

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II. On Mixed Hexanitritonickelates of Praseodymium and Neodymium

the solid phases thereby change to $K_{21}Pr_{5}[Ni(NO_{2})_{6}]g^{H_{2}O}$ and $K_{21}Nd_{5}[Ni(NO_{2})_{6}]g^{H_{2}O}$. There are 10 figures, 8 tables, and 1 Soviet reference.

SUBMITTED:

August 2, 1958

Card 2/2

.5(2)

Tananayev, I. V., Lyutaya, M. D.

TITLE:

AUTHORS:

On the Hexanitrito Nickelates of Samarium, Yttrium, and

Ytterbium (O geksanitranikeleatakh samariya, ittriya 1

itterbiya)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 2,

pp 457-464 (USSR)

ABSTRACT:

The following systems were investigated: $Sm(NO_3)_3 - K_4[Ni(NO_2)_6]$

SOV/78-4-2-31/40

 $H_{2}O$, $Y(NO_{3})_{3}-K_{4}[Ni(NO_{2})_{6}]-H_{2}O$, and $Yb(NO_{3})_{3}-K_{4}[Ni(NO_{2})_{6}]-H_{2}O$

Two solid phases are formed in the first system: $K_5 \text{ Sm}\left[\text{Ni}\left(\text{NO}_2\right)_6\right]_2$ and $K_{19}\text{Sm}_3\left[\text{Ni}\left(\text{NO}_2\right)_6\right]_7 \cdot 4\text{H}_2\text{O}$. The solubility curves suggest the gradual formation of the two phases. The thermograms of the solid phases were plotted and are shown in figures 3 and 4. The thermogram of $K_5 \text{Sm} \left[\text{Ni} \left(\text{NO}_2\right)_6\right]_2$ shows

one endothermal effect only in the temperature range of 225-250°. The thermogram of $K_{19}Sm_3[Ni(NO_2)_6]_7.4H_2O$ shows two

endothermal effects, the first one at 125° and the second one

Card 1/3 in the temperature range of 230-265°C. The compound

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sov/78-4-2-31/40

On the Hexanitrito Nickelates of Samarium, Yttrium, and Ytterbium

 $K_{19}Sm_3[Ni(NO_2)_6]_7.4H_2O$ dissolves more easily in KNO_2 solutions than the respective hexanitrito nickelates of lanthanum, prasoodymium, and neodymium. The compound $K_{19}Y_{3}[Ni(NO_{2})_{6}]_{7}$. $4H_{2}O$ is formed in the system $Y(NO_3)_3-K_4[Ni(NO_2)_6]-H_20$. The thermogram of this compound shows two endothermal effects, the first one at 120° and the second one at 200°-250°. The solubility of $K_{19}Y_3[Ni(NO_2)_6]_7.4H_2O$ in KNO_2 solutions is greater than that of the mixed hexanitrito nickelates of lanthanum, praseodymium, neodymium, and samarium. The hexanitrito nickelate of yttrium is soluble in a 4.6 molar solution of KNO2, whereas the hexanitrito nickelates of praseodymium, neodymium, lanthanum, and samarium are insoluble in this solutions. The phase K_{19} Yo $_3$ [Ni(NO₂)6]7 is formed in the system $Yb(NO_3)_3-K_4[Ni(NO_2)_6]-H_2O.$ The thermographic investigations show an endothermal effect in the temperature range of 190-230°. The thermographic investiations of KNO and

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SOV/78-4-2-31/40 On the Hexanitrito Nickelates of Samarium, Yttrium, and Ytterbium

 ${
m K}_4\left[{
m Ni}\left({
m NO}_2
ight)_6\right]$ were carried out and are shown in figures 9 and 10. The results show that the thermograms of ${
m KNO}_2$ and ${
m K}_4\left[{
m Ni}\left({
m NO}_2
ight)_6\right]$ differ distinctly from all thermograms of the mixed hexanitrito nickelates of rare earths. There are 10 figures, 9 tables, and 3 references, 2 of which are Soviet.

SUBMITTED:

September 15, 1958

Card 3/3

\$/0000/63/000/000/0008/0021

ACCESSION NR: AT4035158

AUTHOR: Samsonov, G. V.; Kosolapova, T. Ya.; Lyutaya, M. D.; Makarenko, G. N.

TITLE: Preparation and physicochemical properties of the carbides and nitrides of the rare-earth elements

SOURCE: AN SSSR. Institut geokhimii i analiticheskoy khimii. Redkozemel'ny*ye elementy* (Rare-earth elements). Moscow, Izd-vo AN SSSR, 1963, 8-21

TOPIC TAGS: rare earth, rare earth element, scandium, lanthanum, yttrium, cerium, carbide, nitride

ABSTRACT: After reviewing the literature on the structure and physical properties (density, melting point, electrical resistivity) of the carbides and nitrides of Sc, Y, La and Ce, the authors describe the preparation of ScC, YC, LaC, ScN, CeN and LaN, the oxidation of the carbides, and some results of an X-ray study of their microstructure. The carbides and nitrides were prepared by heating the oxides with C and N, respectively, at temperatures between 800 and 1800C. The nitrides could also be prepared at lower temperatures by heating the oxide with ammonia. Data are given on the effects of variations in temperature, heating rate and concentration of the reagents, as well as on the relationship between the composition and converged properties of the carbides. Thus, YC2 was found to have the highest

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ACCESSION NR: AT4035158

melting point, electrical resistivity, chemical stability and microhardness, all of which increased with the C/metal ratio. X-ray analysis of the nitrides showed a cubic lattice of the NaCl type with a period of about 4.5-5.5 A. "The X-ray analyses were carried out by 0. T. Khorpyakov." Orig. art. has: 12 figures and 6 tables.

ASSOCIATION: Institut geokhimii i analiticheskoy khimii AN SSSR (Institute of Geochemistry and Analytical Chemistry, AN SSSR)

SUBMITTED: 310ct63

DATE ACQ: 30Apr64

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SUB CODE: IC

NO REF SOV: 016

OTHER: 005

Card ___ 2/2

AUTHOR: Lyutaya, M. D.; Samsonov, C. V.

TITLE: Nitrides of rare dispersed and rare-earth metals

SOURCE: AN Ukr8SR, Institut problem material ovedeniya. Redkiya i redkozemel'ny*ye elementy* v tekhnike (Rare and rare earth elements in engineering). Kiev, Naukova dumka, 1964, 118-126

TOPIC TAGS: Tare metal, rare earth metal, rare metal nitride, rare earth metal/pnitride, gallium nitride, indium nitride, scandium nitride, cerium nitride

ABSTRACT: Nearly pure gallium nitride vas obtained by the treatment of gallium metal (mixed with ammonium carbonate for greater permeation) with nitrogen at 1100C. The nitride obtained resists oxidation bility) with nitrogen at 1100C. It also resists concentrated boiling sulate temperatures up to 700C. It also resists concentrated boiling sulfuric, nitric, and hydrochloric acids, but dissolves completely in furic, nitric, and hydrochloric acids, but dissolves completely in boiling alkall solutions. The reduction of indium seaquioxide with boiling alkall solutions. The reduction of indium seaquioxide with boiling alkall solutions. The reduction of indium seaquioxide with boiling alkall solutions. The reduction of indium nitride decomposes compared to stoichiometric 10.87% nitrogen. Indium nitride decomposes

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n sir at 350C and dissolv cN _{0.97} was obtained by th lack in a nitrogen atmosp ir at temperatures up to com temperature, but decompanthanum and cerium nitrica at 600C. In both case ere obtained. Orig. art.	here. This nitride res 600C. It resists all b imposes in boiling acid des were synthesized by	sists oxidation in out nitric acid at and alkali solutions. Vitreatment with ammo-	0
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ACCESSION NR: AP4041575

S/0078/64/009/007/1529/1533

AUTHOR: Lyutaya, M. D.; Samsonov, G. V.; Khorpyakov, O. T.

TITLE: Germanium nitrides

SOURCE: Zhurnal neorganicheskoy khimii, v. 9, no. 7, 1964, 1529-1533

TOPIC TAGS: germanium nitriding, germanium dioxide nitriding, germanium nitride, germanium nitride structure

ABSTRACT: Conditions of synthesis of germanium nitrides have been studied with 99.99% pure germanium and chemically pure germanium di-oxide as initial materials. Nitriding was performed in ammonia or nitrogen. Germanium nitride with a composition near the stoichiometric composition of Ge₃N₄ was obtained by nitriding in ammonia a mixture of germanium with ammonium carbonate (added to prevent coking) in a 1:2 ratio. Germanium begins to react with nitrogen at 700—750C; at 870C germanium nitride begins to decompose. Nitriding for 1 hr at 800C yielded a nitride with a nitrogen content of 20.52%, compared to the stoichiometric 20.46%. Satisfactory results were also obtained

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with nitriding of germanium dioxide. Addition of ammonium carbonate to germanium dioxide decreased the reaction temperature to 750C and holding time to 1 hr from 800C and 4 hr without ammonium carbonate. X-ray diffraction analysis of the germanium nitride obtained from germanium and germanium dioxide showed that both have rhombohedral structures with the lattice constant a = 8.567Å and α = 107°54'. Germanium nitride is fully resistant to oxidation in air up to 750—800C. In nitrogen it remains stable at temperatures up to 850C. At 900C it decomposes into elements without formation of lower nitrides. Orig. art. has: 2 figures and 6 tables.

ASSOCIATION: Institut metallokeramiki i spetsial'ny*kh splavov AN UkrSSR (Institute of Powder Metallurgy and Special Alloys, AN UkrSSR)

SUBMITTED: 25May63

ATD PRESS: 3065

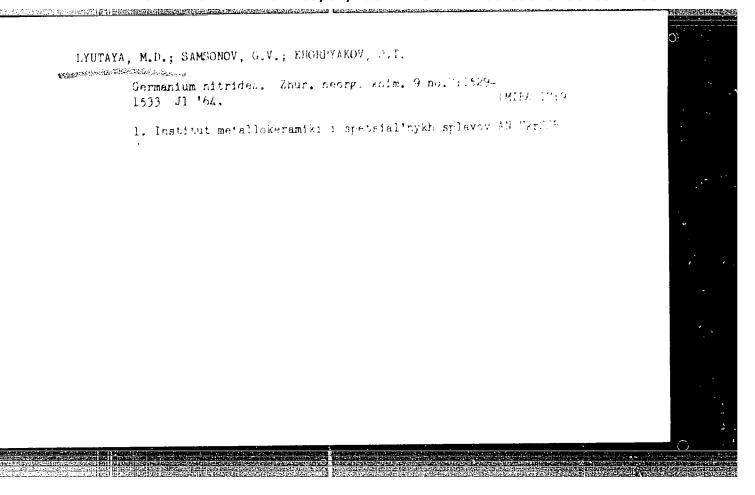
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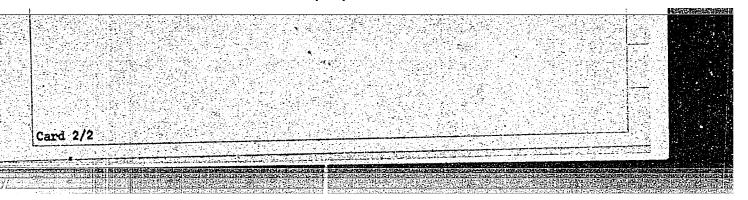
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Card 2/2

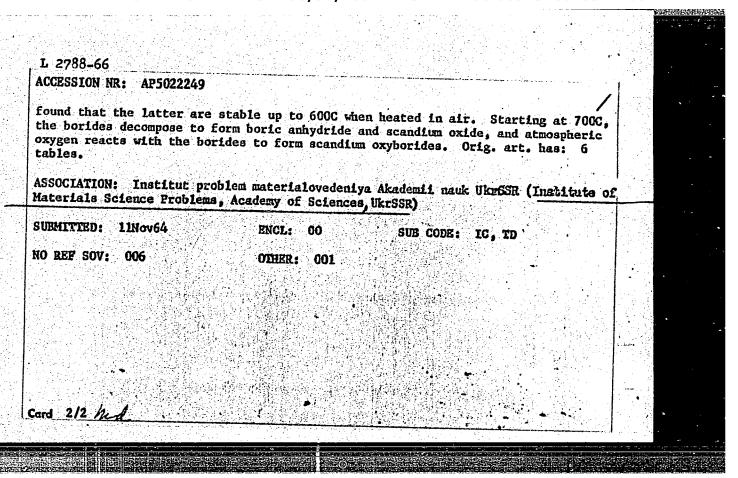


ABSTRACT: Lanthanum germanides (LaGe, LaGe2 and LagGe3) were synthesized in an arc furnace and in a resistance furnace in an argon atmosphere. Composition of the lanthanum germanide product depends upon the ratio of the starting components (elements) and La and Ge vapor pressures. Metallic lanthanum and elemental germanium do not interact up to 900°C. Optimum conditions of preparing lanthanum digermanide (LaGe2) in a resistance furnace are 1 to 1.5 hours melting at 1000-1100°C in an argon atmosphere. Also, reduction of lanthanum oxide with germanium for 1 hour at 1550°C leads to formation of lanthanum digermanide. Lanthanum digermanide dissolves readily in water while LaGe and LagGe3 are practically water insoluble. Thermal

Card 1/2



L 2788-66 EWT(m)/EWP(e)/EWP(i)/ETC/EWG(m)/T/EWP(t)/EWP(b)/EWA(c) IJP(c) JD/JG/AT/WH ACCESSION NR: AP5022249 UR/0363/65/001/007/1039/1043 546.631'271:536.495 AUTHOR: Lyutaya, M. D.; Akinina, Z. S. TITIE: Chemical and thermal stability of scandium borides SOURCE: AN SSSR. Izvestiya. Neorganicheskiye materialy, v. 1, no. 7, 1965, TOPIC TAGS: scandium compound, boron compound, thermal stability, chemical ABSTRACT: Scandium borides ScB₂ and ScB₁₂ were synthesized and their chemical and thermal stability was studied. ScB₂ decomposes in concentrated HCl, H₂SO₄ and HNO3, the decomposition rates being the same in all three acids. ScB12 is stable in HC1 and H2SO4, but decomposes in concentrated HNO3. The effect of dilute acids on both borides is similar to that of concentrated acids. The chemical stability is related to the crystal structure: it increases as the structural elements consisting of boron atoms become more complex. In ScB2, the scandium atoms are insufficiently protected by boron atoms from the action of various reagents; on the contrary, the scandium atoms in ScB12 are well protected by three-dimensional boron networks. In a study of the thermal stability of the scandium borides it is



L 4028-66 EWP(e)/EWT(m)/EWP(t)/EWP(k)/EWP(z)/EWP(b) LJP(c) JD/JG ACCESSION NR: AP5022250 UR/0363/65/001/007/1044/1048 ACCESSION NR: AP5022250 546.76'271:536.495 AUTHOR: Lyutaya, M. Serebryakova, T. I. Thermal stability of chromium borides SOURCE: AN SSSR. Izvestiya. Neorganicheskiye materialy, v. 1, no. 7, 1965, 1044-1048 TOPIC TAGS: chromium compound, boron compound, thermal stability ABSTRACT: The thermal stability of chromium borides Cr4B, Cr2B, Cr3B2, CrB, Cr3B4, and CrB2 in the powdered and compact state was studied in air at 500-1000C. It is found that the borides in the powdered form are practically stable when heated in air up to 600C. The lower borides (Cr2B and Cr3B2) oxidize to form chromium oxyborides and chromic oxide. Cr3B4 and CrB2 decompose on oxidation, forming Cr203 and boric anhydride (B203). The reaction of chromium monoboride with atmospheric oxygen up to 800C forms chromium oxyboride, which decomposes at 9000 to form B203. The most heat-stable borides in the compact state are chromium monoboride and diboride. The great stability of the monoboride up to 900C is due to an oxyboride film which forms on the sample and decomposes at 900-1000C to form B203; the latter serves as the protective film at these Card 1/2

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emperatures. The stabili ormation, upon decomposit roperties in the fused st	ate. Orig. art. has:	high temperatures is due oxide, which has high pro 5 figures and 2 tables.		
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L 11/74-66 EAP(e)/EAT(m)/EAP(1)/EAP(t)/EAP(b) IJP(c) JD/JG

ACCESSION NR: AP5022167

UR/0032/65/031/009/1066/1068 543.77:661.665

AUTHOR: Lyutaya, M. D.; Akinina, Z. S.

74155 Y 11/55 27 27

TITLE: Chemical phase analysis of scandium borides 4455

SOURCE: Zavodskaya laboratoriya, v. 31, no. 9, 1965, 1066-1068

TOPIC TAGS: scandium compound, boron compound, boron, quantitative analysis

ABSTRACT: A method is proposed for determining free boron in ScB₁₂ and ScB₂, based on sintering with barium carbonate. Preliminary experiments show that amorphous boron oxidizes completely to B₂O₃ at 580C and in the presence of BaCO₃ forms a polyborate which is soluble in water. The sinter was treated with water, and boron was determined by titrating with alkali in the presence of mannitol, using phenolphthalein. ScB₁₂ and ScB₂ are stable on heating to 600C in the presence of barium carbonate. Thus, free boron can be determined by sintering the samples with BaCO₃ at temperatures not exceeding 600C. Phase analysis of scandium borides for ScB₂ and ScB₁₂ involved the use of hydrochloric acid solutions, in which ScB₂ decomposes with relative ease, whereas ScB₁₂ remains stable. Results of chemical phase analyses of scandium borides are tabulated. Orig. art.

ACCESSION NR: AP5022167 has: 2 tables. ASSOCIATION: Institut pro Materials Science Problems	blem materialovedeniya . Academy of Sciences U	Akademii naukUko kr89R),,,	SSR(Institute	e of
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32958-66 EWT(m)/EWP(t)/ETI IJP(c) JD/JG ACC NR. AP6015739 (A) SOURCE CODE: UR/0073/66/032/005/0433/0436	
AUTHOR: Lyutaya, M. D.; Goncharuk, A. B.	
ORG: Institute of Problems in the Science of Materials AN UkrSSR (Institut problem materialovedeniya AN UkrSSR)	
TITLE: Chemical properties of lanthanum germanides	•
SOURCE: Ukrainskiy khimicheskiy zhurnal, v. 32, no. 5, 1966, 433-436	
TOPIC TAGS: lanthanum compound, germanium compound, thermal stability, ammonia, analytic chemistry	-
ABSTRACT: The authors study the chemical properties of LaGe ₂ , LaGe and La ₅ Ge ₃ at room temperature in air and in water, and the thermal stability of these compounds in air and in an ammonia atmosphere. The lanthanum germanides used in the study were synthesized from lanthanum and germanium in an arc furnace. Interaction between the german-	
ides and water was determined from the quantity of water passing into solution with accomposition of the lanthanum germanides. Powdered materials with particles measuring the formula stability of the lanthanum germanides. A	
sample of the powder was held at a given temperature for a certain length of time and the oxidation products were then subjected to chemical analysis. Specimens of the three compounds were interacted with ammonia at temperatures of 500-700°, and the prothree compounds	
Card 1/2 UDC: 546.654.289.1	

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L 32958-66

ACC NR: AP6015739

ducts of the reaction were then chemically analyzed. The experimental data are tabulated. It was found that lanthanum digermanide LaGe $_2$ is easily decomposed by moisture while LaGe and La $_5$ Ge $_3$ are practically stable in water. There is a direct relationship between the composition of the lanthanum germanides and their stability in air and in an ammonia atmosphere. Lanthanum digermanide is more resistant to atmospheric oxidation at 200-400° and to interaction with ammonia at 500-700° than are LaGe and La $_5$ Ge $_3$ under the same conditions. Intermediate compounds, lanthanum hydroxygermanides, are formed in the germanide oxidation process. A compound with the empirical formula LaGeN is produced by interactions between LaGe and ammonia at 500-700°. La $_5$ Ge $_3$ also interacts with ammonia in this temperature range. Orig. art. has: 1 figure, 4 tables.

SUB CODE: 07/ SUBM DATE: 09Nov64/ ORIG REF: 005/ OTH REF: 002

Card 2/2

L 44361-66 EWT(m)/EWP(k)/EWP(e)/EWP(t)/ETI IJP(c) JD	
ACC NR: AP6007295 SOURCE CODE: UR/0226/66/000/002/0108/0109	
REPORTER: Lyutaya, M. D.; Goncharuk, A. B.	
ORG: none	
TITLE: All-Union Inter-Institute Seminar on the Production, Properties and Applications of the Nitrides (held in Kiev from 20 to 22 April 1965)	
SOURCE: Poroshkovaya metallurgiya, no. 2, 1966, 108-109	
TOPIC TAGS: metallurgic conference, nitride, nitride compound, metallurgic research	
ABSTRACT: The Seminar was attended by ~100 delegates from >30 research and academic institutions of the Soviet Union. 32 papers were presented. They dealt with such topics as: a classification of nitrides based on theories of their electron structure and chemical bonding (G. V. Samsonov); production of the nitrides of rare-earth, rare, disseminated and transition	
metals (M. D. Lyutaya and others); experimental production of aluminum mitride from the gaseous phase (N. G. Slavina and A. A. Pletyushkin); production of transition-metal hitrides by nitriding metal powders and reducing and nitriding metal oxides (G. V. Samsonov and V. S.	
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"APPROVED FOR RELEASE: 08/31/2001

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L 44361-66 10 ACC NR: AP6007295 Polishchuk); research into nitride alloys (L. M. Katanov and others); fabrication of work parts from nitrides (L. I. Struk and others); research into the evaporation of nitrides (V. V. Fesenko and others); studies of the superconductivity, and thermoemissive and refractory properties of the nitrides (O. I. Shulishova and others). S. M. Ariya and associates presented an interesting paper on the formational enthalpy of titanium nitrides as a function of their composition, while T. N. Nazarchuk presented a general survey of methods of the chemical analysis of nitrides. The resolution adopted by the Seminar noted the high level of the presented papers and outlined further ways and means of enhancing the effectiveness of nitride research. In particular, it pointed to the need to intensify R&D work on high-purity nitrides and to broaden the studies of the physical properties of nitrides and nitride-base alloys by utilizing x-ray spectral, galvanomagnetic, magnetic, spectroscopic and other methods. The proceedings of the Seminar will be published in a special volume. SUB CODE: 11_20_3 07/ SUBM DATE: none/ 2/2

ACC NRI AP6020961

SOURCE CODE: UR '0226/66/000/006/0060/0063

AUTHOR: Lyutaya, M. D.; Goncharuk, A. B.

ORG: Institute for Problems in Science of Materials, AN UkrSSR (Institut problem materialovedeniya AN USSR)

TITLE: Lanthanum germanides

SOURCE: Poroshkovaya metallurgiya, no. 6, 1966, 60-63

TOPIC TAGS: germanide, lanthanum, lanthanum germanide, germanothermic method germanide. Sturyesis, CHEMICAL REQUERION

ABSTRACT: The authors have investigated conditions for synthesizing lanthamum germanides LaGe₂, LaGe, and La₅Ge₃ from elements in an arc furnace and lanthamum digermanide by the germanothermic method, i.e., reduction of lanthamum oxide with germanium in vacuo. Some chemical properties of lanthamum germanides,

Card 1/2

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ACC NR: AP6020961

stability of germanide powders in water, and thermal stability in air and in ammonia have been studied. It has been shown that the thermal oxidation of lanthanum germanides in air is followed by the formation of lanthanum hydroxygermanides as wintermediate products. The interaction of lanthanum monogermanide with ammonia yields an LaGeN product. Orig. art. has: 2 figures and 2 tables. [Based on authors abstract]

SUB CODE: 11/ SUBM DATE: 18Mar66/ ORIG REF: 003/ OTH REF: 002/

Cord 2/2

SHTROBEL', V.; ROMANKOV, P.G.; KONOVALOV, V.I.; LYUTAYA, N.S.

Study of mass transfer in a rotor-disk extractor. Zhur.prikl.khim.
(MIRA 17:2)

1. Leningradskiy tekhnologicheskiy institut imeni Lensoveta.

SHTROBEL', V.; ROMANKOV, P.G.; KONOVALOV, V.I.; LYUTAYA, N.E.

Study of hydrodynamics without mass transfer and in the presence of mass transfer in a rotor-disk extractor. Zhur. prikl. khim.
36 no.12:2672-26RO D'63. (MIRA 17:2)

1. Leningradskiy tekhnologicheskiy institut imeni Lensoveta.

SOYFER, V.M.; LYUTAYA, V.A.

Using a silica composition for the rammed lining of small steelpouring ladles. Ognsupory 30 no.10:5-6 '65. (MIRA 18:10)

1. Khar'kovskiy zavod "Elektrotyazhmash" im. V.I. Lenina.

LYUTAYFVA, V. A.

Lyutayeva, V. A. -- "Paraff'notherapy of exceeds ive them'is," Sborn'k trudov (Tomskiy obl. nauch.-issled. in-t fiz. metodov lecheniya i kurortologii), Vol. VI, 1949, p. 183-88

S0: U-5241, 17 December 1753, (Letonis 'zhurnal 'nykh Statey, Ho. 26, 1949).

TAROSLAVSKIY, M.I.; LYUTENBERG, R.M.; CHERNYSHOV, V.N.

Instrument for the analysis of the piezoelectric properties of crystals. Zhur.tekh.fiz.26:499-441 F '56. (MIRA 9:6)

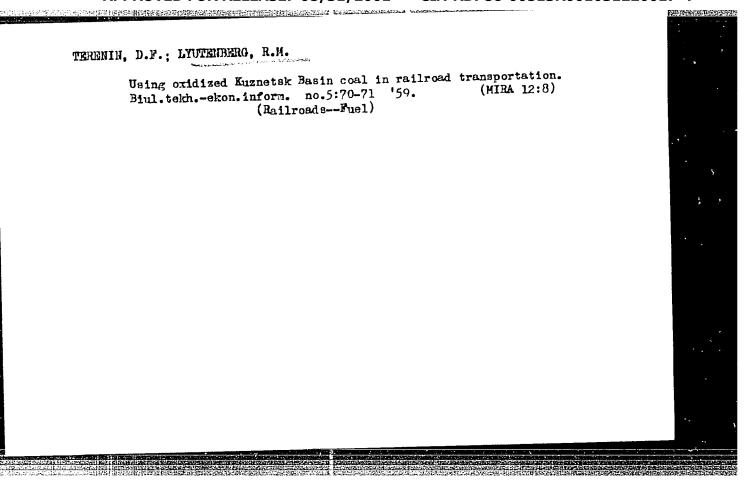
(Piezoelectricity--Heasurement)

TERENIN, D.F., kand, tekhn, nauk; LYUTENBERG, R.M., inzh.

Quality of oxidized coals of the Kuznetek Basin and their use in locomotives. Vest. TSNII MPS 16 no.8:24-30 D '52.

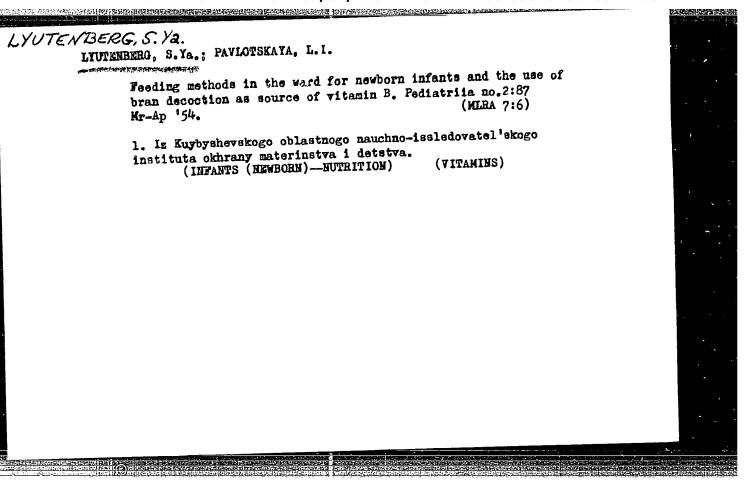
(MIRA 11:1)

(Locomotives) (Kuznetek Basin--Coal--Analysis)



KORCHEV, V.V., kand, tekhn.neux; LYUTENBERG, R.M., inzh.

Testing of diesel fuels in a small cylinder capacity engine. Vest.
TSHII MPS 22 no.9:27-30 '63. (MIRA 17:2)



LYUTENKO, AN

AUTHOR:

None Given

72-2-19/20

TITLE:

For the Industry of Ceramics - a Progressive Technology (Keramicheskoy promyshlennosti - peredovuyu tekhnologiyu).

PERIODICAL:

Steklo i Keramika, 1958.

Nr 2, pp. 46-47 (USSR)

ABSTRACT:

A technical conference of the functionaries of the ceramic industry took place in Khar'kov in December 1957, which was organized by the Ukrainian administration of the Scientific-Technical Society of the building material industry and the Ministry of Building Material Industry of the Ukrainian SSR. The conference was attended by functionaries of the works producing ceramics in the Ukraine and the Russian Federation, the Economic Councils of Stalinsk and Khar'kov, the state-controlled offices for Economic Planning of the USSR, the RSFSR, and the Ukrainian SSR, the Building- and Building-Material Department of the TsK KPU and of the Scientific Research- and Planning Institutes. The results obtained in the Ukrainian Ceramic Industry and prospects for the future were discussed. Particular attention was paid to the utilization of progressive experience in the industry as well as to the introduction of new technical methods,

Card 1/4

high-efficiency equipment, and a progressive technology.

For the Industry of Ceramics - a Progressive Technology

72-2-19/20

- 1.) I.I.Moroz (Minister for the Building Material Industry of the Ukrainian SSR) delivered a report on the work and the prospects of the ceramics industry.
- 2.) A.A.Kopeykin (Director of the NIIstroykeramiki) spoke about the work carried out by his institute. He was reproached for talking too much about future plans and too little about work already completed.
- 3.) A.A.Grebennik (Head of the PKB NIIstroykeramiki), after his report, was criticized for the same reasons as Kopeykin.
- 4.) Dudnik (TsKB MPSM Ukrainian SSR, Khar'kov) spoke about the introduction of new equipment and assembly lines.
- 5.) N.I.Dikerman (Chief Engineer of the Administration of the Mosstroymaterialy) stated that the efficacy of the brick charging devices for tunnel kilns at present no longer corresponds to the increased efficiency of the kilns.
- 6.) A.N.Lyutenko (Chief Engineer of the Administration of the Economic Council, Khar'kov) spoke about production reserves of plants.
- 7.) S.M.Beluga (Chief Engineer of the Metlakh Tile Works, Khar' kov) spoke about the mechanization of production.

Card 2/4

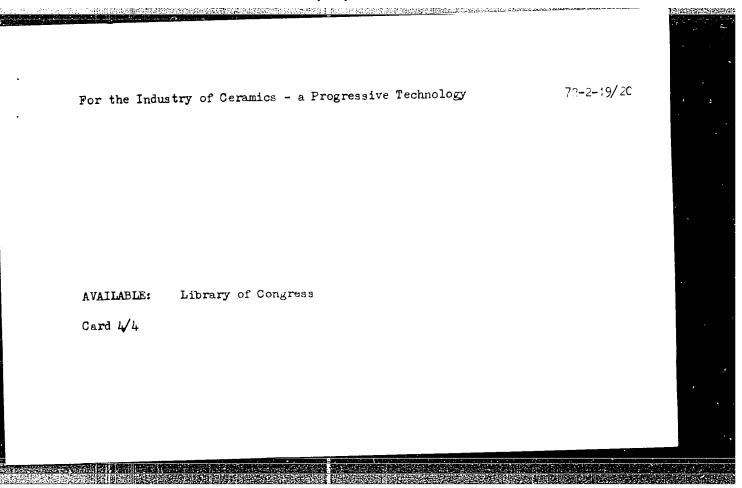
For the Industry of Ceramics - a Progressive Technology

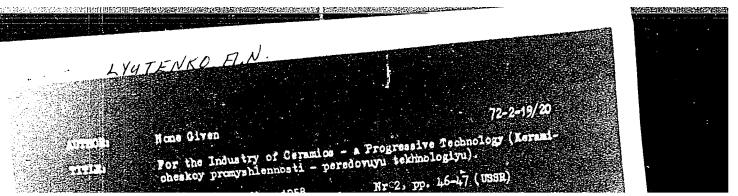
72-2-19/20

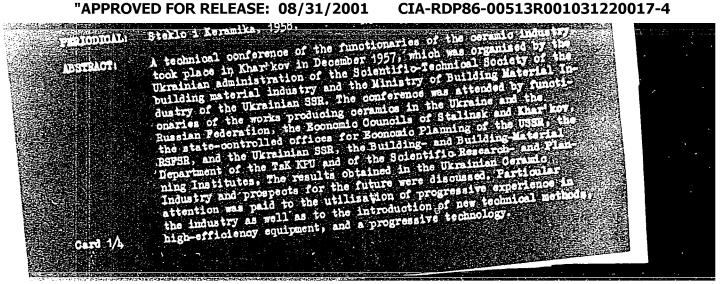
- 8.) L.K.Parnovskiy (Director of the Ceramics Factory, Lvov) spoke about success achieved in production.
- 9.) P.Ye.Andrianov delivered a report on the ceramics industry of Italy.
- 10.) M.D.Abramovich (Director of the Combined Plant "Keramik" at Kiyev) spoke about the organization of the production of mosaic tiles.
- 11.) S.M.Brekhovskikh (Chief Specialist for Glass of the Gosplan USSR) criticized the lack of reports concerning the stage of furnace technology.
- 12.) A.N.Lyutenko, G.A.Soldatov, S.M.Beluga, M.V.Gordyga and F.K.Perre reported on the unfavorable situation of the raw material sector, which impairs the delivery of high-quality raw materials to factories and plants.

Decisions were made for the purpose of improving industrial work, for the purpose of reducing time needed for smelting and drying, with a view of speeding up mechanization and improving the quality of products, as well as of increasing production and reducing initial costs.

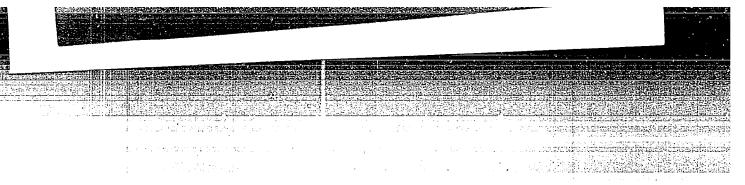
Card 3/4

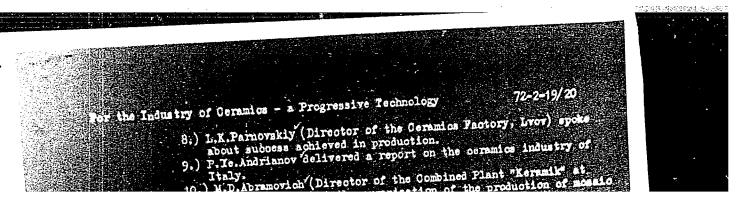


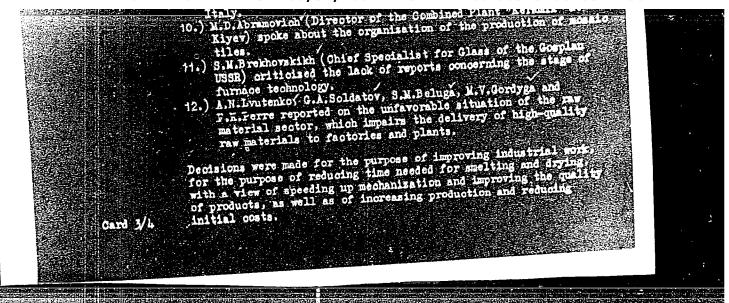




APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"







AUTHOR:

Lyutenko, A Ye

SOV-128-58-10-15/19

TITLE:

Experience in the Application of Cherkassy Bentonite in the Production of Steel Casting (Opyt primeneniya cherkasakogo Bentonita v proizvodatve stal nogo lit ya)

PERIODICAL:

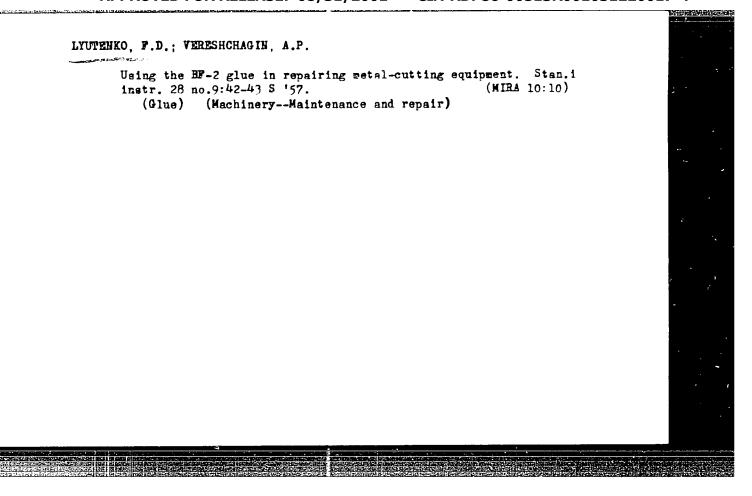
Liternoye proizvodstvo, 1958, Nr 10, p 31 (UDDR)

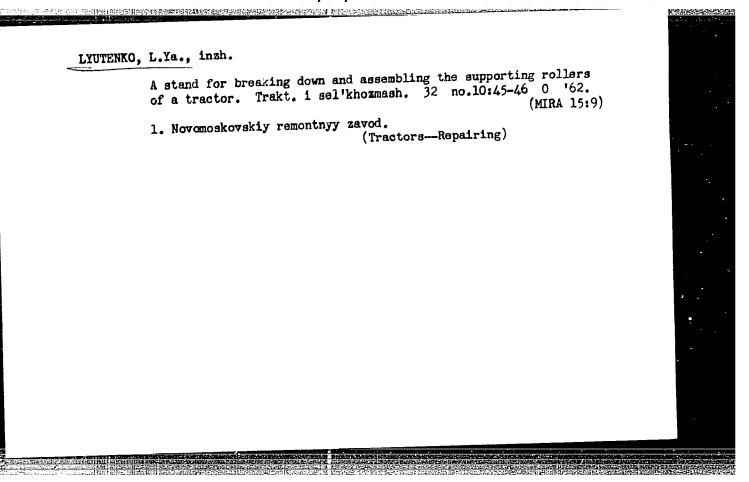
ABSTRAIT.

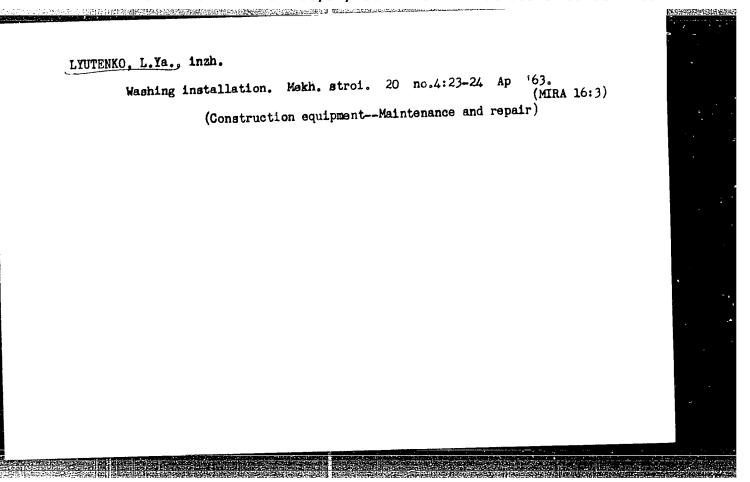
The zavod 'Irpen mashtorf' ("Irpen mashtorf" Flant; used lotal 40/70 sand in mold-making for pig iron and steel castings. Removal of the sand crust picked up in casting was a negative factor. The Institut Mashinovedeniya (Institute of Nechanical Engineering; under the direction of Corresponding Member of AS Tkr SSF, A A. Jorshkov, recommended the use of bentonite from the Markasay (Ukraine) region. The author conducted production experiments in the plant, and obtained good results with this bentonite in mold and core making. Since then the plant has used this bentonite for 14 months with positive results. Mixture percentages are given; and it is recommended that bentonite be used in the foundries of the Kiyev Conomic Region.

1. Steel castings--Production 2. Bentonite--Applications

Card 1/1







APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

L 15691-65 FSF(h)/FSS-2/EWT(1)/EEC(m)/FS(v)-3/EWG(s)-2/EWG(v)/FCC/EWA(d)/ EEC-4/EEC(t)/EWA(h) Po-4/Pe-5/Pq-4/Pg-4/P1-4/P1-4/Pac-2/Peb/Pb-4 AEDC/ AFFTC/AFMDC/ESD-3/RADC/AFGC/ESD(t)/ESD(E1)/AEDC(a)/SSD/BSD/AFWJ/AFMDC/AFETR/ ACCESSION NR: AP5000175 AFTC(b)/AFTC(a)/ASD-3 \$/0293/64/002/006/0928/0932

AUTHOR: Shafer, Yu. G.; Sokolov, V. D.; Skryabin, N. G.; Lyutenko, V. F.; Yary*gin, A. V.; Salimzibarov, R. B.

TITLE: Intensity distribution of cosmic rays in the atmosphere to a height of 500 km

SOURCE: Kosmicheskiye issledovaniya, v. 2, no. 6, 1964, 928-932

TOPIC TAGS: solar activity cycle, cosmic ray, geophysical rocket, single counter, ionization camera, Kosmos satellite, cosmic ray albedo, magnetic storm

ABSTRACT: In the period from 1958 to 1963, during a decrease in solar activity, cosmic ray measurements have been carried out by means of geophysical rockets and satellites of the Kosmos type. Geophysical rockets were equipped with single counters and ionization cameras. Satellites of the Cosmos type were equipped with ionization cameras, single counters, and counting telescoped for measuring the cosmic ray albedo. Rocket and satellite launchings were scheduled for days without magnetic storms and quiet sun. Primary cosmic rays were measured at heights of 100--500 km. The cosmic ray albedo measured by rockets equipped with special

Card 1/2

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

L 15691-65

ACCESSION NR: AP5000175

devices was found to be insignificant. Numerical values of measurement data show a slight increase in particle count with height. No indications were found which would associate systematic variations in the intensity of primary commic rays with the eleven-year cycle of solar activity. Orig. art. has: 1 figure and 3 tables.

ASSOCIATION: none

SUBMITTED: 13May64 ENCL: 00 SUB CODE: AA, SV.

NO REF SOV: 003 OTHER: 008 ATD PRESS: 3144

LYUTENKO, V.F., inzhener.

Correcting the frequency of a quartz crystal. Vest.sviazi 16 (MIRA 10.10)

1. Yakutskiy radiotsentr. (Oscillators, Crystal)

LIMITANC, V.F., inzh.; ZAMYATIN, K.M., tekhnik.

LYUTENKO, V.F., inzh.; ZAMYATIN, K.M., tekhnik.

Using a heterodyne wavemeter as an exciter. Vest.sviazi 17 no.8:

(MIRA 10:10)

32-33 Ag *57.

1.Yakutskiy radiotsentr (for Lyutenko).

(Radio--Transmitters and transmission)

\$/845/62/000/004/001/013 E192/E382

9. 5.

AUTHORS: Grigorov, N.L., Sokolov, V.D. and Lyutenko, V.F.

TITLE: Measurement of slow neutrons from cosmic rays on

aircraft

SOURCE: Akademiya nauk SSSR. Yakutskiy filial. Trudy. Seriya

fizicheskaya. no. 4. 1962. Variatsii intensivnosti

kosmicheskikh luchey, 4 - 9

TEXT: An instrument for measuring slow neutrons in cosmic rays is described. This is based on the principle of an integrating ionization chamber which is filled with gaseous BF₃.

The ionization current of such a device consists of two components: that due to the decay of B and that due to all the remaining ionizing effects. 10 In order to separate the ionization current due to the decay of B the current produced by the other effects is compensated by a second ionization chamber. The wo chambers are spherical with internal diameters of 20 cm; the walls are steel, 1 mm thick. The collector electrodes are connected together into one electrical system. The capacitance of the chambers with the system of electrodes is about 12 pF. The spheres are electrically Card 1/3

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5/845/62/000/004/001/013 E192/E382

Measurement of

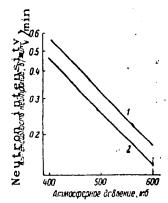
insulated and voltages of opposite signs are applied to them. A certain charge produced by the neutron-ionizing current is stored on the collecting electrode of the chambers during a unit time Δt ; the magnitude of this charge is proportional to the intensity of the neutrons and the charge can be measured by the method described by N.I. Grigorov (UFN, 8, no. 4, 1956). In this method the charge on the collector electrode is converted into a voltage pulse of definite magnitude. The pulses so obtained are applied to the input of a athode-follower tube; the pulse is negative and is of about 850 us duration. This pulse is applied to an amplifier and then to a nonlinear amplifying stage, where it is lengthened to about 40 msec but where its amplitude is still proportional to that of the input pulse. It is then fed to a switching audio circuit, whose output signal is in the form of an audio pulse of 3 kc/s; the duration of this audio pulse is proportional to the charge stored on the collector electrodes of the chambers. The audio pulse is applied to a counter which records the number of cycles. The circuit for measuring the charge is based on directly-heated tubes. The equipment was used between August 24 - 29, 1959, in flying Card 2/3

S/845/62/000/004/001/013 E192/E382

Measurement of

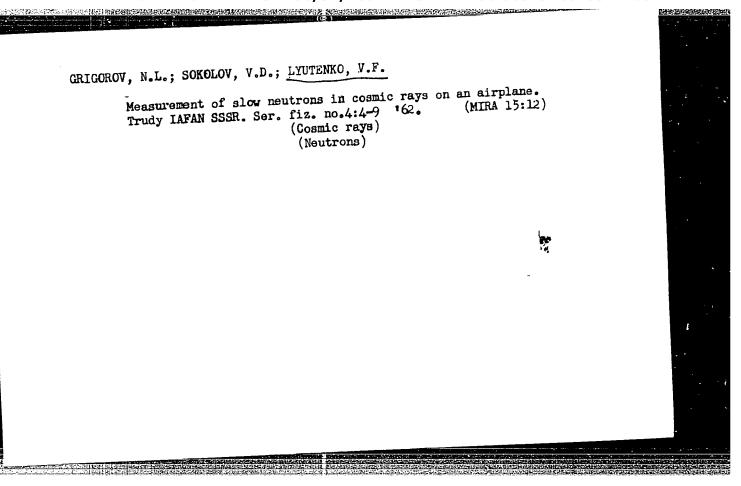
aircraft at Yakutsk. It was first checked by using ${\rm Co}^{60}$ γ -source. The neutron intensity was measured as a function of the atmospheric pressure and the results averaged over five flights are illustrated in Fig. 3. There are 3 figures.

Fig. 3:



Atmospheric pressure, mb

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KAZANSKIY, B.A.; DOROGOCHINSKIY, A.Z.; ROZENGART, M.I.; LYUTER, A.V.;
MITROFANOV, M.G.

Aromatization of marrow hexane fractions of Groznyi gasoline on an alumina-chromic oxide catalyst. Kin.i kat. 1 no.2:201-209
(MIRA 13:8),
Jl-Ag '60.

1. Institut organicheskoy khim:i im. N.D.Zelinskogo AN SSSR i Groznenskiy nauchno-issledovatel skiy neftyanoy institut.
(Aromatization)
(Hexane)

5.3300

Dorogochinskiy, A. Z., Lavrent'yev, V. I.,

s/020/60/131/02/045/071 B011/B011

Lyuter, A. V., Mel'nikova, N. P.,

Kupriyanov, V. A.

68998

TITLE:

AUTHORS:

Synthesis and Properties of Naphthenic Hydrocarbons With a Long

Side Chain

PERIODICAL:

Doklady Akademii nauk SSSR, 1960, Vol 131, Nr 2, pp 367 - 370

(USSR)

ABSTRACT:

The authors wanted to work out a general method and conditions for the synthesis of technical fractions of the substances mentioned in the title, as well as the study of the properties of these fractions. Propylene, butylene, amylene, hexylene, and heptylene were used for the purpose. As a result of the experiments conducted at the authors institute, a 3-stage scheme of synthesis was suggested: 1) synthesis of olefins with a given number of C-atoms, or polymerization, respectively. A dehydrated pentane-amylene fraction from thermal cracking, purified from the sulphur compounds, was utilized. The catalyst was phosphoric acid on kieselgur. Olefins with ramified structure were obtained in this connection. The highest yield of isodecenes occurred at 170-180°, pressure of 50-60 atm, volume rate 3-4 h-1. Amylenes

Card 1/4

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4" Synthesis and Properties of Naphthenic Hydrocarbons With a Long Side Chain

68998 S/020/60/131/02/045/071 B011/B011

were transformed to 70%. A concentrate boiling between 120 and 185° was obtained from the polymerizate (yield 85-90%). Table 1 shows the resulting (mostly ramified) structures of isodecenes. Table 2 shows their physico-chemical properties (the raw material was fraction 6 of the thermal cracking and benzene). Isomerization and hydro-dehydro polymerization of the olefins were ascertained as side reactions. 2nd stage: alkylation. Aromatic hydrocarbons (benzene, toluene) were alkylated by means of the isodecenes produced (Refs 3-5). The best conditions were: 97% H2SO4, reaction time 2 hours, ratio benzene:isodecene = 5:1. Temperature 10-20°. The alkylate amounted to 140% by weight of olefins or 90% of the theoretical yield. A fraction boiling between 180° and 350° was obtained from the alkylate as a concentrate of isodecyl benzenes (85% of the alkylate). It chiefly consisted of mono-substituted derivatives of benzene (Table 2). On using aluminum chloride as catalyst the yield was higher and attained 97-98%. Disproportionation occurred as side reaction. 3rd stage: hydrogenation. The alkylate concentrate was hydrogenated on 2 catalysts: a) nickel catalyst. The optimum conditions were: pressure 7 atm, molar ratio hydrogen:alkylate = 2.8:1; 150-2000.

Card 2/4

Synthesis and Properties of Naphthenic Hydrocarbons S/020/60/131/02/045/071 With a Long Side Chain 68998

Synthesis and Properties of Naphthenic Hydrocarbons B011/B011

Volume rate 0.2 h 1; b) nickel-tungsten catalyst. Optimum conditions: pressure in the reaction zone 200 atm; molar ratio hydro-gen-alkylate = 64:1; 300°; volume rate 0.5 h⁻¹. To prevent a temperature increase on the latter catalyst, the alkylate was diluted with gasoline distillate (fraction 80-120°) of the trade-mark "Kalosha" in a ratio of 1:2. Destruction was recorded as a side reaction. The desired naphthene fraction was obtained from the hydrogenation product by rectification. It boils out between 180° and 350°. Its yield attained 90% of the aromatic hydrocarbons contained in the alkylate (Table 2). The range of the fluctuation of properties in dependence on procedure and raw materials is shown in table 3. Data obtained show that the scheme described here leads to naphthene hydrocarbons with a long side chain, high density, high calcricity, and a low freezing temperature. The following names are mentioned: Ye. G. Vol'pova, L. A. Potolovskiy, I. F. Blagovidov, L. I. Kostikin, Yu. A. Gol'dshtein, Yu. I. Kozorezov, A. Z. Dorogochinskiy, and K. I. Zimina. There are 3 tables and 6 Soviet references.

Card 3/4

68999

Synthesis and Properties of Naphthenic Hydrocarbons S/020/60/131/02/045/071 With a Long Side Chain 8011/8011

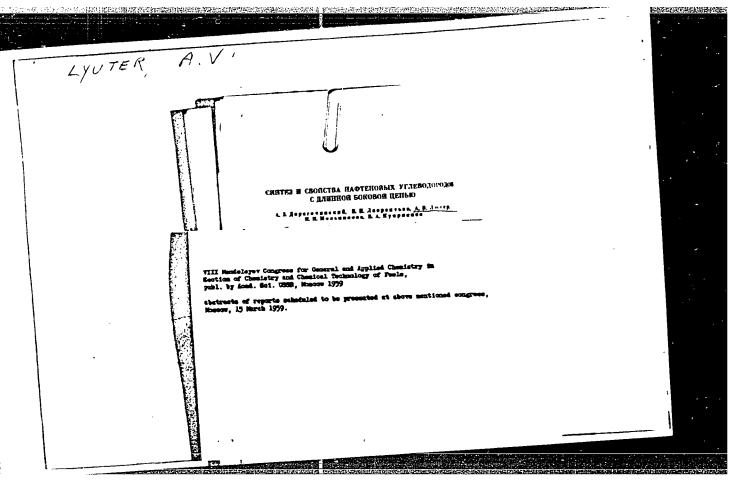
ASSOCIATION: Groznenskiy neftyanoy nauchno-issledovatel'skiy institut

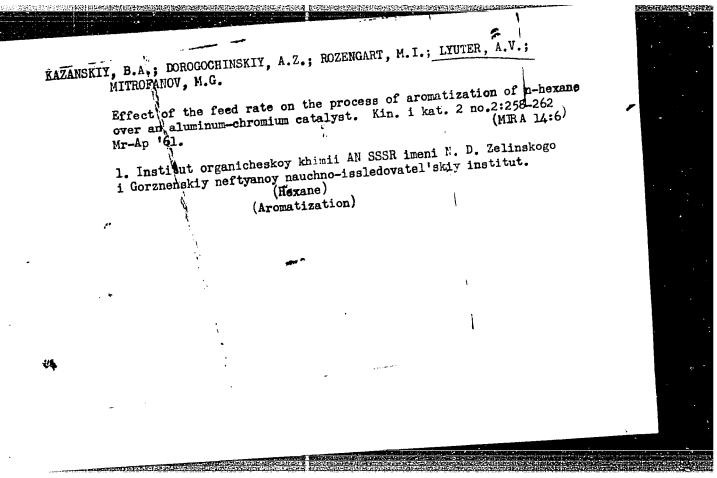
(Groznyy Scientific Research Institute of Petroleum)

PRESENTED: November 28, 1959, by B. A. Kazanskiy, Academician

SUBMITTED: November 25, 1959

Card 4/4





APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

CIA-RDP86-00513R001031220017-4 "APPROVED FOR RELEASE: 08/31/2001 To be the little and the second of the second secon

5/204/62/002/004/003/019 E071/E433

AUTHORS:

Kazanskiy, B.A., Dorogochinskiy, A.Z., Sterligov, O.D.,

Lyuter, A.V., Dmitriyevskiy, M.L., Nazarov, P.S.

TITLE:

Dehydrogenation of isopentane into isoamylenes on an

alumochromopotassium catalyst

PERIODICAL: Neftekhimiya, v.2, no.4, 1962, 448-456

A systematic study of the process of dehydrogenation of isopentane into isoamylenes under conditions of a stationary and moving layer of granulated catalyst K-544 was carried out on experimental installations of Groz NII. Tests on the stationary layer were carried out on a laboratory and an enlarged installation. The reactors with a stationary layer of the catalyst were of the capacity of 40 and 500 cm3 respectively. Tests in the moving layer were made in a co-current continuous. pilot plant with a reactor (4 litres) and a regenerator (4.7 litres). The volume of the catalyst - 35 litres, throughput - about 100 litres/day, the velocity of circulation of the catalyst up to 16 litres/hour. The analyses of the reaction products were made by chromatographic and other chemical methods. The influence of the temperature, volume velocity and rate of recirculation of Card 1/2

> APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

s/204/62/002/004/003/019 E071/E433

Dehydrogenation of isopentane ...

the catalyst on the main parameters of the process as well as the behaviour of the catalyst were studied. It was found that the catalyst had a good and stable activity. During an operating period of 1100 hours in a stationary layer and 400 hours in a moving layer its activity remained practically unchanged. the optimum condition of the process (temperature - 540°C and volume velocity - 1 hour-1) the yield of isoamylenes amounted to 30 to 31 wt.% calculated on raw material (98.6% of isopentane) with a total yield of unsaturated hydrocarbons C5 of 34 to 38 wt.%. The catalyst has a satisfactory strength and good The velocity of burning out of coke from the most inaccessible layers of catalyst K-544 amounted to 20 litres/litre of catalyst per hour, in comparison with that for aluminosilicate catalysts of 13 to 16 litres/litre of There are 6 figures and 5 tables. catalyst per hour.

ASSOCIATION: Institut organicheskoy khimii AN SSSR im. N.D.Zelinskogo (The Institute of Organic Chemistry AS USSR imeni N.D.Zelinskiy) GrozNII

Card 2/2

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

KAZANSKIY, B.A.; DORCCCCHINSKIY, A.Z.; ROZENGART, M.I.; TYUN'KINA, N.I.;

KUZNETSOVA, I.M.; LXUTER, A.V.; MITROFANOV, M.T.

Aromatization of mixtures of n. hexane with 2-methylpentane,
with 3-methylpentane or methylcyclopentane. Izv.AN SSSR.Otd.
(NIRA 15:7)
khim.nauk no.7:1308-1309 Jl '62.

1. Institut organicheskoy khimii im. N.D.Zelinskogo AN SSSR.
(Aromatization) (Paraffins)

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

DOROGOCHINSKIY, Akiviy Zinov'yevich; LYUTER, Aleksandr Valentinovich;
VOL'POVA, Yevgeniya Grigor'yevna; REKHVIASHVILI, Antonina
Nikolayevna; ROLESNIKOV, F.M., red.; KUZ'MENKOVA, N.T.,
tekhn. red.

[Oil gases in the Chechen-Ingush and other economic regions
of the Northern Caucasus]Neftianye gazy Checheno-Ingushakogo
i drugikh ekonomicheskikh raionov Severnogo Kawkaza. Groznyi
i drugikh ekonomicheskikh raionov Severnogo (MIRA 16:3)

(Caucasus, Northern—Gas, Natural)

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

KAZANSKIY, B.A.; DOROGOCHINSKIY, A.Z.; ROZENGART, M.I.; GITIS, K.M.;

LYUTER, A.V.; MITROFANOV, M.G.

Effect of the length of an alumine-chromic-potassium
catalyst layer on the aromatization of n-heptane.

Kin.i kat. 4 no.2:315-318 Mr-Ap '63. (MIRA 16'5)

1. Institut organicheskoy khimii AN SSSR imeni N.D.Zelinskogo i
Groznenskiy neftyanoy nauchno-issledovatel'skiy institut.

(Heptane) (Aromatization) (Catalysts)

KAZANSKIY, B.A.; DOROGOCHINSKIY, A.Z.; S.ERLIGGY, O.D.; LYUTER, A.V.;

IMITRIYEVSKIY, M.L.; NAZAROVA, M.P.; REMIVIASHVILI, A.N.

Studying the dehydrogenation of isopentane on K-544 and K-5

finely divided catalysts. Trudy GrozNII no. 15:241-253 '63.

(MIRA 17:5)

KAZANSKIY, B.A.; DOROGOCHINSKIY, A.Z.; ROZENGART, M.I.; LYUTER, A.V.;
MITROFAMOV, M.G.; BRESHCHENKO, Ye.M.; KALITA, L.A.; GOL'DSHTEYN,
Yu.A.; AFANAS'YEV, A.I.; MAYAR'YEV, S.V.; ZAMARV, V.V.

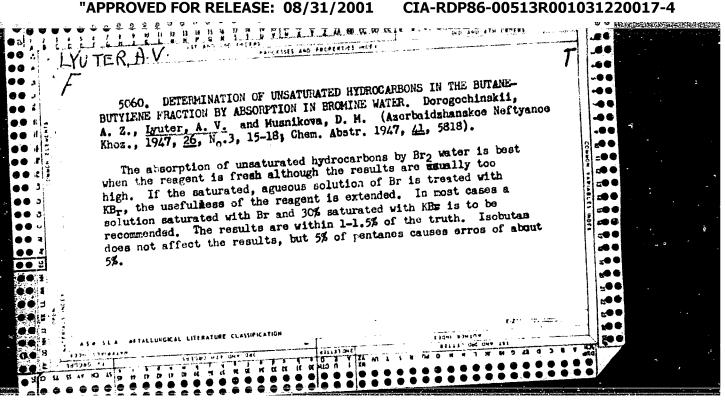
Dehydrocyclization of normal hexane. Trudy GrozNII no. 15:
254-264 '63.

(MIRA 16:5)

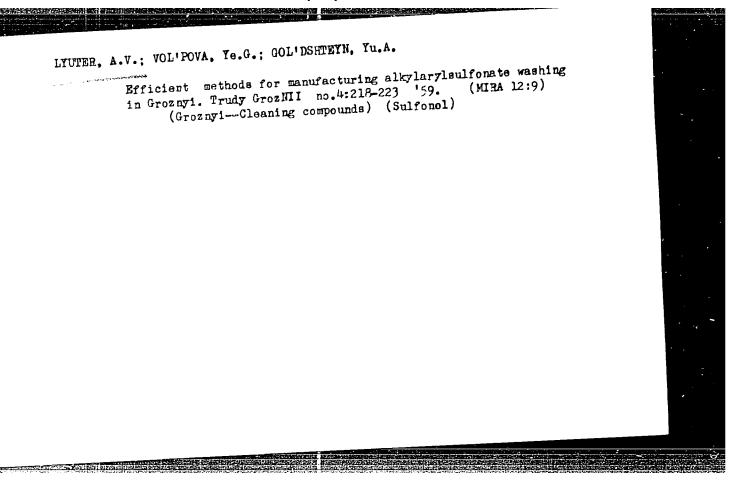
KAZANSKIY, B.A.; DOROGOCHINSKIY, A.Z.; ROZENGART, M.I.; KUZNETSOVA, Z.F.; LYUTER, A.V.; MITROFANOV, M.G.

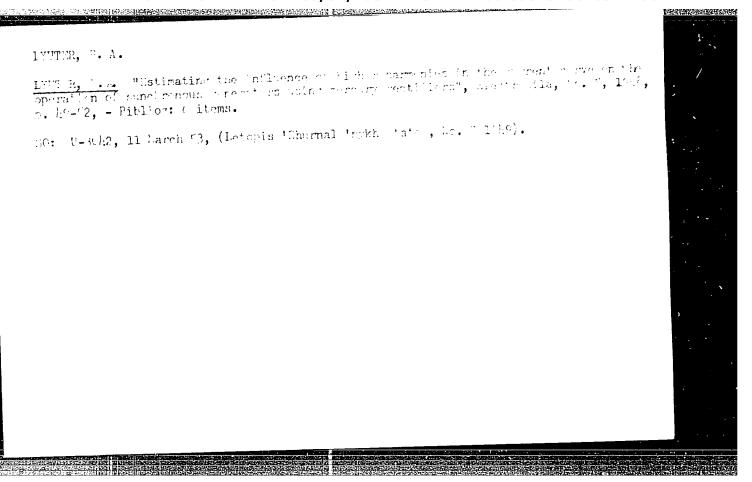
Changes in alumina-chromia catalysts during the aromatization of n-hexane. Kin.i kat. 4 no.5:768-772 S-0 '63. (MIRA 16:12)

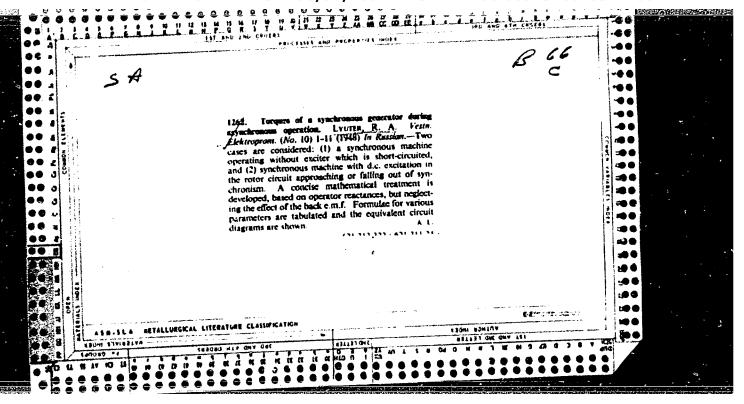
1. Institut organicheskoy knimii AN SSSR imeni N.D.Zelinskogo i Groznenskiy neftyanoy nauchno-issledovatel'skiy institut.



APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"







LYUTER. A. A.

Nov 48

USSR/Elecricity - Machinery, Design Currents, Electric Direct

"Physical Limitations in DC Fachines and Certain Related Problems of Design," V. 1. Kas'yanov, A. A. Kashin, R. A. Lyuter, I. N. Rabinovich, D. V. Shapiro, Laureates of Stalin Prize, Elektrosil' Factory imein Stalin, 5 pp

"Vest Elektro-Prom" No 11

Points out increasing importance of DC machines and new design problems cropping up. Devotes attention to problem of increasing the ultimate power of DC motors for a given diameter of their armatures—pe mitting an increase in motor speed, decrease in weight and dimensions, and decrease in flywheel moment for reversible motors. Analyzes means of increasing ultimate power and commutiting ability, and improving design, and points out other problems requiring attention.

PA 65/49T42

ORLOVSKIY, A.V., professor; LYUTER, R.A., doktor tekhnicheskikh nauk; KAZOVSKIY, Ye.Ya., kandidat tekhnicheskikh nauk; YAKOBSON, El'mar, inzhener; ANTOPOL'-SKIY, V.M., inzhener; PUKHOV, G.Ye., doktor tekhnicheskikh nauk; FYUHSTEN-BERIN, A.I., inzhener; BERGER, A.Ya., professor (Leningrad); TSVERAVA, G.K., inzhener; KRAYNIY, K.I., inzhener (g.Kotovsk, Tambovskoy obl.); BELOV, V.N., inzhener (g.Ul'yanovsk).

Correspondence conference of readers of "Elektrichestvo" Elektrichestvo no.8:89-91 Ag 153. (MLA 6:8)

1. Kiyevskiy politekhnicheskiy institut (for Orlovskiy). 2. Zavod "Elektrosila" (for Lyuter and Kazovskiy). 3. Estonkommunenergo (for Yakobson).
4. Saratovskiy industrial'nyy tekhnikum (for Antopol'skiy). 5. Tomskiy politekhnicheskiy institut imeni Kirova (for Pukhov). 6. Tikhvinskiy glinozemnyy zavod (for TSverava). (Electric engineering—Periodicals)

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

AID P - 3024

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Card 1/2

Subject : USSR/Electricity

Pub. 27 - 11/33 Authors

: Alekseyev, A. Ye., Corr. Memb. Academy of Sci. Prof. of USSR, A. S. Yeremeyev, Eng., and R. A. Lyuter, Dr. of Tech. Sci.

: Problems of the domestic water-wheel generator design Title

Periodical: Elektrichestvo, 7, 55-65, Jl 1955

: The tremendous development of hydroelectric power stations in the USSR creates the problem of designing Abstract

more and more powerful water-wheel generators. This

in turn places several technical problems to be solved by Soviet machine manufacturers and designers. Among these problems are: rationalized grouping of all the elements; static and dynamic stability under operation for long transmission lines; improvements in construction details, like that concerning the total height of the generators in relation to the rotation speed of the water wheel, etc. Eleven

APPROVED FOR RELEASE: 08/31/2001 CIA-RDP86-00513R001031220017-4"

AID P - 3024

Elektrichestvo, 7, 55-65, J1 1955

Pub. 27 - 11/33 Card 2/2

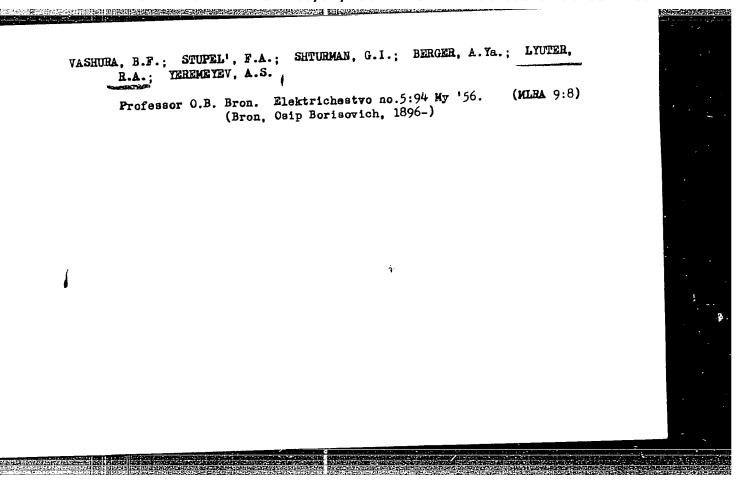
photographs, drawings and diagrams, 3 tables, 15 Soviet references (1945-1955) 1 American (1934).

Institution: Plant "Elektrosila" im. Kirov.

Submitted: Ap 13, 1955

ALL VERDYANTS, L.M.; KLETSKIKH, I.N.; KOSTENKO, M.P.; LYUTER, R.A.;
SAPOHNIKOV, R.A.; CHAPLINSKIY, S.K.; CHEMERKIN, K.G.

I. V. Tokov; obituary. Elektrichestvo no.12:77 D '55. (MLRA 9:3)
(Tokov, Ivan Vasil'evich, 1901-1955)



IVANOV, N.P.; YEHEMEYEV, A.S.; LYUTER, R.A.; KAPLAN, M.Ya.; IPATOV, P.M.

Powerful hydrogenerators. Elektrosila no.14:5-11 '56.
(Hydroelectric power stations)

(Hydroelectric power stations)

Dynamoelectric excitation system for hydrogenerators. Elektrosila no.14:33-35 '56. (MIRA 12:12) (Electric generators)

8(0) SOV/112-59-1-698

Translation from: Referativnyy zhurnal. Elektrotekhnika, 1959, Nr 1, pp 92-93 (USSR)

AUTHOR: Lyuter, R. A., Samoylovich, N. Ya., and Koldobskiy, M. I.

TITLE: Thermal Durability of Squirrel-Cage-Rotor AC Electric Machinery

PERIODICAL: Elektrosila, Nr 15, 1957, pp 29-42

ABSTRACT: Heating of induction and synchronous motors is examined under these conditions: starting, undervoltage operation, cutting-off one phase of the synchronous motor, overload up to the limit of steady-state stability, and excitation loss. Temperature rise in ${}^{O}C$ of the starting rotor winding during the starting period is $\Theta_{C} = C$; $(1 - e^{-t_{\rm n}/T_{\rm s}})$, where w is the average value of losses during starting per unit surface of bars in w/cm²; C = 20-100 degrees cm²/w is the heating constant of piece bars over the steel (it depends on the tightness of bar-steel contact), roughly $C \approx 50$; $T_{\rm s}$ is the time constant of bar heating for round copper bars; with C = 50, $T_{\rm s} \approx 44$ d_C per sec, where d_C is

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the bar diameter in cm; $t_n = \frac{T_m M_H}{M_n K_u}$ is the starting time in seconds.

' (Translator's note: Apparently, the first formula is incorrectly typeset in the Russian original.) The quantity of heat evolved in the rotor over the starting period with the initial slip s of the rotating rotor is

$$Q_p = \frac{s^2}{2} T_m M_H \frac{1}{K_u}$$
 in kw. sec, where

$$T_{m} = \frac{27.4 \text{ GD}^2 (n_{N}/100)^2}{M_{H}}$$
 is the mechanical time constant in sec;

$$\frac{1}{K_u} \approx \frac{1}{1-M_c/M_n}; \ M_H \ \text{is the rated motor torque in synchronous kw; n_w is} \\ m_n \ \text{and } M_c \ \text{are the starting torque and the resistance torque of the drive (in the largest content of the drive of the drive)}$$

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synchronous kw), both being functions of the slip s in the general case; GD² is the flywheel effect of all spinning masses in ton · m². In simplified calculations, under the assumption of adiabatic heating, the temperature rise over the starting period of the rotor starting winding made from copper, brass, or bronze can be computed from the formula $\Theta = 1.28 \frac{t_n M_n}{G} k_k k_b$ in °C where G is the starting winding weight in kg; the coefficients $k_k = 0.80$ -0.90 and $k_b = 1$ for a single-cage winding; $k_k k_b = 0.60$ -0.75 for a double-cage motor whose upper cage weighs G. Assuming one hot starting and two cold startings with the rotor temperature rise of $\frac{\Theta_{max}}{k_k k_b} = 250$ °C for single-cage induction motors and 300°C for synchronous and double-cage induction motors, the maximum starting time permissible by rotor heating conditions will be $t_{n-max} = 195 \frac{G}{M_n}$ for single-cage induction motors and

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 $t_{n \text{ max}} * 235 \frac{G}{M_{n}}$ for synchronous and double-cage induction motors. On the

basis of stator heating conditions, assuming a temperature rise of 35-40°C per one starting for class-A insulation windings and of 50-55°C for class-B insulation windings, the permissible starting time in seconds will be

 $t_{n}^{\dagger} \max = \frac{7,850}{j_{nH}^2}$ for class-A insulated windings and $t_{n}^{\dagger} \max = \frac{10,500}{j_{nH}^2}$ for

class-B insulated windings, where j_{DH} is the initial starting current density in amp/mm². Estimated values of permissible starting time are between 4 and 15 sec. In 3-kv synchronous and induction motors, the starting time is limited by rotor overheating, while in 6-kv induction motors, by stator overheating. With an undervoltage and motor operation within its stable range, the permissible time of operation with the voltage 1 - p as a fraction of the rated

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Thermal Durability of Squirrel-Cage-Rotor AC Electric Machinery

voltage is
$$t_p^1 = \frac{1.25}{I_p^{1/2} - 1} t_{1.5}$$
 in seconds, where $t_{1.5}$ is the standard

permitted 50%-current overload time (GOST 183-55 specifies 60 and 120 sec); the stator current in induction machines I_p^i as a fraction of the rated current is determined, for undervoltage conditions, from the current diagram for the specified active power; in the synchronous machines the field current, as a fraction of the rated current, for undervoltage conditions, should be determined from the vector diagram for the field current. In case of a considerable

undervoltage, the deceleration time of the motor is $T^1 = T_m \frac{M_H}{M_c - M_n/(1 - p^2 cdk)}$

Over the time required to attain the slip s the rotor-winding temperature rise

will be
$$\Theta'_{s} = 1.28 \frac{T' M_{Tx} (1 - p)^{2} s^{2}}{G} k_{k} k_{b} \text{ in } {}^{O}C.$$

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Thermal Durability of Squirrel-Cage-Rotor AC Electric Machinery

The permissible speed drop, for undervoltage conditions, can be determined from this rule: over the deceleration time down to the slip s and over the subsequent speed-rise time on voltage recovery, the rotor-winding temperature rise should not exceed the specified value Θ_{\max} in ^{O}C . Hence

$$s = \sqrt{\frac{\Theta_{\text{max}} M - C. \text{ Nehce}}{1.28 [T' M_n (1 - p)^2 + T_{\text{rn}} M_h / k_u] k_k k_b}}.$$

When the motor is operating with one phase cutoff, its stator current is equal to the line-to-line voltage divided by the sum of positive-phase-sequence and negative-phase-sequence impedances. The time of one-phase-off operation is largely determined by heating the rotor with negative-phase-sequence currents

$$I_2(b q/e)$$
. The quantity $A_2 = \int_0^t I_2^2 dt$, where t in seconds should not exceed

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120-150 for induction motors, about 60 for synchronous motors (except for 2-pole types), and about 30 for 2-pole synchronous motors. Permissible time of under-load operation of a synchronous motor on loss of field can be determined in a way similar to the undervoltage case, i.e., considering the value of stator or rotor current and the value of $t_{1.5}$.

Ye. Ya. K.

Card 7/7

AUTHORS:

Kostenko, M.P., Alekseyev, A. 10., 30V105-58-7-50/52

Lyuter, R.A., Zavalishin, D. A. Gnedin, L. F., Britsin, N. L.

TIT .:

Leonid Nikolayevich Gruzov (Deceased)

PERIODICAL:

Elektrichestvo, 1958, Hr 7, pp 93-93 (USSH)

ABJTRACT:

Professor Leonid Nikolayevich Gruzov, Dector of Technical Sciences, Engineer-Colonel, Head of the Kafedra elektropitaniya ustanovok svyazi Voyennov krasnoznamennov akademii svyazi

ustanovok svyazi Voyennoy krasnoznamennoy akademii svyazi (Department of Electric Supply of Telecommunication Equipment at the Krasnoznamennaya Military Academy of Telecommunication) a prominent expert in the field of electric machines, died on October 17th, 1957, at the age of 51. He graduated with distinction from the Donskoy politekhnicheskiy institut (Don Polytechnical Institute) in 1927, was then aspirant at the Leningradskiy politek nicheskiy institut (Leningrad Polytechnical Institute), assistant, and finally docent at the same institute. He combined his scientific and pedagogical activity with that of an engineer. He published a series of papers on the transient modes of operations of electric machines and of power supply systems.

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THE REPORT OF THE PARTY OF THE

Leonid Aikolayevich Gruzov

30V/105-58-7-30/32

He worked as engineer in the "Elektrosila" works as well. He took part in Morld War II. In 1947 he took his degree as Doctor of Technical Sciences. He developed a method for the investigation of electric machines. He was first head of the Department of Theoretical Electrical Engineering, then of the Department of Electric Supply Plants at the Military Academy of Telecommunication. He published nore than 30 scientific papers, textbooks and manuals. There is 1 photograph.

1. Scientific personnel--USSR

Card 2/2

ZHERVE. Georgiy Konstantinovich; LYUTER, R.A., doktor tekhn.nauk, retsenzent; RIVLIN, L.B., inzh., red.; SOBOLEVA, Ye.M., tekhn.red.

[Industrial testing of electric machinery] Promyshlennye ispytaniia elektricheskikh mashin. Izd.2., perer. Moskva, Gos.energ.izd-vo, 1959. 504 p.

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IVANOV, V.I.; LIUTER, R.A.; MANOYLOV, V.Ye.; YERMOLIN, N.P.;
FRAMKE, A.V.

Vladimir Tikhonovich Kas'ianov; on the seventy-fifth anniversary
of his birth and the tenth anniversary of his death.
Elektrichestvo no.4:95 Ap '62. (MIRA 15:5)

(Kas'ianov, Vladimir Tikhonovich, 1887-1952)

IPATOV, P.M., kand.tekhn.nauk. KAZOVSKIY, Ye.Ya., doktor tekhn.nauk;
KULIKOV, N.V., inzh.; LIUTEK R.A., doktor tekhn.nauk;
Research conducted at the Leningrad branch of the All-Union
Scientific Research Institute of Electromechanics and the S.M.
Kirov "Elektrosila" factory. Vest.elektrprom. 33 no.413.8

Ap '62. (Electric machinery)

(Electric machinery)

KOSTENKO, M.P., akademik; LYUTER, R.A., doktor bekhr.nauk; KAZOTEKIT, Tele, doktor tekhn.sauk, prof.; IVANOV, N.P., kand.tekhr.nauk

Conditions governing the use of nonsynonronous cutting-is in electric power systems. Elektrichestvo no.12:77-78 D 65.

(MIRA 18:12)